

COATED FUEL CELL ELECTRODES

CROSS-REFERENCE TO RELATED APPLICATIONS

This is a Continuation-in-Part of application Serial Number 09/795,930, filed February 28, 2001, which is based on Provisional Serial Number 60/185,924.

BACKGROUND OF THE INVENTION

Electrodes for proton exchange membrane fuel cells are typically made by mixing a slurry of carbon black with deposited platinum particles together with a liquid dispersion of ion conducting polymer solution. This slurry is then cast into a film and dried and the resulting film is a porous electrode that has a three-dimensional reaction zone with ionic conducting paths, electronic conducting paths and void spaces for reactant or byproduct transport. A continuing need for such structures is to balance the water content of the electrode and the underlying membrane with the open void space of the electrode for oxygen transport. It is known that the oxygen diffusion rate in water is roughly three to four orders of magnitude less than that in free air. Thus, if the catalyst sites are blocked with a film of water with any significant thickness, the reaction rate slows due to lack of oxygen transport. On the other hand, if not enough water is incorporated in the ionomer part of the electrode, then the ionic conductivity of the ionomer is drastically reduced, and the overall resistance of the cell increases. This balance is referred to in the literature as "water management."

SUMMARY OF THE INVENTION

The instant invention provides electrodes which aid in water management in fuel cells.

Specifically, the instant invention provides an electrode suitable for use in a fuel cell bearing a coating of at least one transport polymer. The invention further provides, in a fuel cell comprising at least one cathode and at least one anode and an ion exchange membrane separating the cathode and the anode, the improvement wherein at least one electrode bears a coating of at least one transport polymer.

BRIEF DESCRIPTION OF THE DRAWING

Figure 1 is a schematic, cross-sectional illustration of a fuel cell of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

A wide variety of coating polymers can be used in the present invention. When used on the anode, the coating functions as an oxygen transport polymer. When used on the cathode, it functions as a hydrogen transport polymer. The basic requirement of

aid, the functioning of an electrolytic cell. In practice, these oils are believed to prevent the ionomer components in the cathode electrode from drying out, while continuing to permit electrons and oxygen to pass. In addition, the oils exclude water from the voids of the cathode electrode layer, substituting a material that has an affinity for oxygen. Moreover, the beneficial effect is long lasting, in that the oxygen or hydrogen transport polymers typically have a low vapor pressure, which results in insignificant loss due to vaporization.

An additional advantage of applying a coating to an electrode of a fuel cell is the mitigation of carbon monoxide poisoning on the anode catalyst when operating on reformed hydrocarbon fuel streams. The transport mechanisms of most coatings previously investigated rely on the principle of a molecular sieve. Thus the coating material will more readily pass small molecules and will block the transport of large molecules. Since the CO molecule is more than ten times larger than the hydrogen molecule, the application of a similar coating to the anode catalyst site would reduce the exposure of that catalyst to the CO.

Typically in PEM fuel cell systems, the CO content of the fuel stream needs to be reduced to 50 ppm or less. The present invention provides significant fuel processing benefits, in that the coating reduces the amount of CO that reaches the catalyst, thereby reducing the need for CO removal.

The present invention is further illustrated by the following specific example.

EXAMPLE:

A catalyst layer coating was made by dispersing 40 wt% Pt on carbon available from Englehard in NAFION™ solution. The catalyst solution was coated onto a decal material and dried under forced hot air. The resulting electrode was hot pressed at 135°C for 3 minutes onto a 1 mil thick NAFION™ ionomer membrane. The cathode side electrode was coated with perfluoropolyether (PFPE) solution and dabbed “dry” with an adsorbent wipe. Weighing the sample before and after application of the PFPE solution indicated that 18 mg of the PFPE was applied over 25 cm². The sample was tested in a fuel cell at varying temperatures and humidification conditions and found to have performance superior in all cases to control samples made at the same time which were identical except for the PFPE treatment.